

Pills * * * For Skin Purification, * * * For Pallor, Weakness and Nervousness, * * * For Sick Headache, Indigestion * * * Digestive * * * For * * * Chills and Grip."

On December 20, 1930, the Potter Drug & Chemical Corporation, Malden, Mass., having withdrawn its claim and answer and consented to the entry of a decree, judgment of condemnation and forfeiture was entered, and it was ordered by the court that the product be destroyed by the United States marshal.

ARTHUR M. HYDE, *Secretary of Agriculture.*

17859. Misbranding of Amogen tablets. U. S. v. 3 Dozen Bottles of Amogen tablets. Default decree of condemnation, forfeiture, and destruction. (F. & D. No. 25311. I. S. No. 713. S. No. 3565.)

Examination of samples of a drug product, known as Amogen tablets, from the herein-described interstate shipment having shown that the labels bore claims of curative and therapeutic properties that the article did not possess, the Secretary of Agriculture reported the matter to the United States attorney for the Southern District of California.

On November 13, 1930, the United States attorney filed in the District Court of the United States a libel praying seizure and condemnation of three dozen bottles of Amogen tablets, remaining in the original unbroken packages at Los Angeles, Calif., consigned by the Amogen Co., San Antonio, Tex., alleging that the article had been shipped from San Antonio, Tex., on or about October 11, 1930, and transported from the State of Texas into the State of California, and charging misbranding in violation of the food and drugs act as amended.

Analysis of a sample of the article by this department showed that it consisted essentially of calomel and extracts of plant drugs, including a laxative drug and a mydriatic drug.

It was alleged in substance in the libel that the article was misbranded in that the following statements regarding the curative or therapeutic effects of the said article, and similar statements in Spanish, appearing on the bottle label and in the accompanying circular, were false and fraudulent: (Bottle) "Indigestion * * * for the Liver;" (circular) "Headache * * * Indigestion * * * Influenza, La Grippe * * * Kidney and Liver Troubles, Malaria Conditions, Sores in Mouth, Loss of Appetite."

On December 15, 1930, no claimant having appeared for the property, judgment of condemnation and forfeiture was entered, and it was ordered by the court that the product be destroyed by the United States marshal.

ARTHUR M. HYDE, *Secretary of Agriculture.*

17860. Adulteration and misbranding of elixir potassium bromide, tincture nux vomica, tincture digitalis, sodium salicylate tablets, and phenolphthalein tablets. U. S. v. Brewer & Co. Plea of guilty. Fine, \$500. (F. & D. No. 25006. I. S. Nos. 02430, 02431, 02435, 02559, 05741.)

Examination of the herein-described drugs showed the following results: The elixir potassium bromide contained less potassium bromide than required by the National Formulary; the tincture nux vomica contained more of the alkaloids of nux vomica than the maximum prescribed by the United States Pharmacopoeia; the tincture digitalis had a lower potency than required by the pharmacopoeia; and the sodium salicylate tablets and the phenolphthalein tablets contained smaller amounts of the respective drugs than declared on the labels.

On September 30, 1930, the United States attorney for the District of Massachusetts, acting upon a report by the Secretary of Agriculture, filed in the District Court of the United States for the district aforesaid an information against Brewer & Co. (Inc.), a corporation, Worcester, Mass., alleging shipments by said company in violation of the food and drugs act, from the State of Massachusetts into the State of Maine, on or about August 25, 1928, of a quantity of tincture nux vomica; on or about September 5, 1928, of a quantity of elixir potassium bromide and sodium salicylate tablets; on or about October 19, 1928, of a quantity of phenolphthalein tablets; and on or about November 13, 1928, of a quantity of tincture digitalis. which said drugs were adulterated and misbranded. The articles were labeled in part as set out below.

Adulteration of the elixir potassium bromide was alleged for the reason that it was sold under and by a name recognized in the National Formulary.